

United States Patent and Trademark Office

UNITED STATES DEPARTMENT OF COMMERCE United States Patent and Trademark Office Address: COMMISSIONER FOR PATENTS P.O. Box 1450 Alexandria, Virginia 22313-1450 www.usplo.gov

APPLICATION NO.	FILING DATE	FIRST NAMED INVENTOR	ATTORNEY DOCKET NO.	TORNEY DOCKET NO. CONFIRMATION NO	
10/789,266	02/26/2004	Thomas Jessel	5199-169	9559	
Leslie Gladston	7590 04/30/2007 e Restaino, Esq.	EXAM	EXAMINER		
Brown Raysman Millstein Felder & Steiner LLP			GAMETT, DANIEL C		
163 Madison Avenue P.O. Box 1989		ART UNIT	PAPER NUMBER		
Morristown, NJ 07962-1989			1647		
			MAIL DATE	DELIVERY MODE	
			04/30/2007	PAPER	

Please find below and/or attached an Office communication concerning this application or proceeding.

The time period for reply, if any, is set in the attached communication.

Office Action Summary		Application	No.	Applicant(s)					
		10/789,266		JESSEL ET AL.					
		Examiner		Art Unit					
		Daniel C. Ga	·	1647					
Period fo	The MAILING DATE of this communication app or Reply	ears on the co	over sheet with the co	orrespondence address					
WHIC - Exter after - If NO - Failu Any	ORTENED STATUTORY PERIOD FOR REPLY CHEVER IS LONGER, FROM THE MAILING DATE in time may be available under the provisions of 37 CFR 1.13 SIX (6) MONTHS from the mailing date of this communication. It is period for reply is specified above, the maximum statutory period were to reply within the set or extended period for reply will, by statute, reply received by the Office later than three months after the mailing and patent term adjustment. See 37 CFR 1.704(b).	ATE OF THIS 36(a). In no event, will apply and will ex, cause the applicate	COMMUNICATION however, may a reply be tim topice SIX (6) MONTHS from topic to become ABANDONED	l. ely filed he mailing date of this communication. D (35 U.S.C. § 133).					
Status									
1)⊠	Responsive to communication(s) filed on 14 Fe	ebruary 2007.							
2a) <u></u> □	This action is FINAL . 2b)⊠ This action is non-final.								
3)	Since this application is in condition for allowance except for formal matters, prosecution as to the merits is								
	closed in accordance with the practice under Ex parte Quayle, 1935 C.D. 11, 453 O.G. 213.								
Dispositi	on of Claims								
5) <u></u> 6)⊠	Claim(s) 1-67 is/are pending in the application. 4a) Of the above claim(s) 1-42 and 53-67 is/are Claim(s) is/are allowed. Claim(s) 43-52 is/are rejected. Claim(s) is/are objected to. Claim(s) are subject to restriction and/or	e withdrawn fr			•				
Applicati	ion Papers								
10)🖾	The specification is objected to by the Examine The drawing(s) filed on <u>26 February 2004</u> is/are Applicant may not request that any objection to the Replacement drawing sheet(s) including the correct The oath or declaration is objected to by the Ex	e: a)⊠ accep drawing(s) be h ion is required	neld in abeyance. See if the drawing(s) is obj	37 CFR 1.85(a). ected to. See 37 CFR 1.121(d).				
Priority ເ	under 35 U.S.C. § 119								
 12) Acknowledgment is made of a claim for foreign priority under 35 U.S.C. § 119(a)-(d) or (f). a) All b) Some * c) None of: 1. Certified copies of the priority documents have been received. 2. Certified copies of the priority documents have been received in Application No. 3. Copies of the certified copies of the priority documents have been received in this National Stage application from the International Bureau (PCT Rule 17.2(a)). * See the attached detailed Office action for a list of the certified copies not received. 									
Attach	**(a)								
2) Notice 3) Information	ct(s) ce of References Cited (PTO-892) ce of Draftsperson's Patent Drawing Review (PTO-948) ce of Draftsperson's Patent Drawing Review (PTO-948) ce No(s)/Mail Date 01/18/2005 12/13/2006	5)	Interview Summary Paper No(s)/Mail Da Notice of Informal P	te					

Application/Control Number: 10/789,266 Page 2

Art Unit: 1647

DETAILED ACTION

1. Applicant's election of claims 43-52 in the reply filed on 02/14/2007 is acknowledged. Because applicant did not distinctly and specifically point out the supposed errors in the restriction requirement, the election has been treated as an election without traverse (MPEP § 818.03(a)).

- 2. Applicant's election of FGF as the species of modulator of the FGF signaling pathway in the reply filed on 02/14/2007 is acknowledged. Applicant expressed some confusion as to the requirement for election of species. Applicant's assumption that the intent was to identify claim 49, and select from activators of an FGF signaling pathway is correct, especially in view of section 7 of the requirement for election/restriction mailed on 08/14/2006. Because applicant did not distinctly and specifically point out the supposed errors in the restriction requirement, the election has been treated as an election without traverse (MPEP § 818.03(a)).
- 3. Claims 1-42 and 53-67 are withdrawn from further consideration pursuant to 37 CFR 1.142(b) as being drawn to a nonelected inventions, there being no allowable generic or linking claim. Election was made without traverse in the reply filed on 02/14/2007.
- 4. Claims 43-52 are under consideration insofar as the read upon methods wherein the activator of an FGF signaling pathway is an FGF protein.

Claim Rejections - 35 USC § 112

5. The following is a quotation of the first paragraph of 35 U.S.C. 112:

The specification shall contain a written description of the invention, and of the manner and process of making and using it, in such full, clear, concise, and exact terms as to enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to make and use the same and shall set forth the best mode contemplated by the inventor of carrying out his invention.

Application/Control Number: 10/789,266

Art Unit: 1647

6. Claims 43-52 are rejected under 35 U.S.C. 112, first paragraph, because the specification, while being enabling for methods for identifying a modulator of an FGF signaling pathway wherein the activator of an FGF signaling pathway is an FGF protein, does not reasonably provide enablement for methods wherein the activator of an FGF signaling pathway is anything other that an FGF protein. The specification does not enable any person skilled in the art to which it pertains, or with which it is most nearly connected, to use the invention commensurate in scope with these claims. Claims 43-49, 51 and 52 are drawn to methods for identifying a modulator of an FGF signaling pathway. The methods comprise incubating cells with an "activator of an FGF signaling pathway". The specification provides at [0176-0177] a broad definition and non-limiting exemplary list for the term "activator of an FGF signaling pathway". The list appropriately includes each of the known FGF proteins. The also list includes molecules which are not known to be directly involved with FGF signaling and for which any association with FGF signaling stems from common usage of intracellular pathways (e.g. ras, PKC, PI3K) shared by many growth factors. The list therefore includes signaling molecules that are not specific to FGF signaling, some of which are identified in the specification. VEGF, for example, is identified as an activator of both Wnt and FGF signaling [0143, 0177]. TGFβ is identified as an activator of BMP, Hh, and FGF signaling [0154, 0163, 0177]. The specification provides no guidance as one could identify a modulator of an FGF signaling pathway if any of these molecules were used as the "activator of FGF signaling" in the claimed method. If, for example, VEGF were the "activator of FGF signaling", a test compound that gave a measurable result would identified as a modulator of VEGF signaling, but the result would not show that the test

Application/Control Number: 10/789,266

Art Unit: 1647

Page 4

compound acts on a component of VEGF signaling shared with FGF. Further experimentation would be required to determine if the test compound actually modulates an FGF signaling pathway. The same can be said of any of the named growth factors and cytokines in [0177] and claim 49. The claimed method cannot identify a modulator of an FGF signaling pathway unless a known activator of an FGF signaling pathway is used. By encouraging the skilled artisan to employ molecules that are only coincidently related to FGF signaling, the specification actually guides the skilled artisan away from identifying a modulator of an FGF signaling pathway.

7. Claim 50 is directed to a method of identifying a modulator of neural differentiation. The recited steps could achieve this goal because they do provide for a determination of neural differentiation. However, this claim, along with claims 43-49, 51 and 52, still directs the artisan to contact cells with any one of a plethora of "activators". Many of the "activators" are intracellular molecules, such as ras, raf Grb2, PI3 kinase, MEK, PIP2, PKC, PLC, PTN, and etc. The specification does not provide for cells that do not intrinsically possess these molecules and so does not teach the skilled artisan how to controllably contact cells with these "activators". In addition to the multitude of named "activators", the list also includes "any analogue or homologue thereof". The skilled artisan would be required to perform undue experimentation order to practice any of the methods of claims 43-52 with this indeterminably large genus of reagents.

Application/Control Number: 10/789,266

Art Unit: 1647

Claim Rejections - 35 USC § 102

Page 5

8. The following is a quotation of the appropriate paragraphs of 35 U.S.C. 102 that form the

basis for the rejections under this section made in this Office action:

A person shall be entitled to a patent unless -

(b) the invention was patented or described in a printed publication in this or a foreign country or in public use or on sale in this country, more than one year prior to the date of application for patent in the United States.

(e) the invention was described in (1) an application for patent, published under section 122(b), by another filed in the United States before the invention by the applicant for patent or (2) a patent granted on an application for patent by another filed in the United States before the invention by the applicant for patent, except that an international application filed under the treaty defined in section 351(a) shall have the effects for purposes of this subsection of an application filed in the United States only if the international application designated the United States and was published under Article 21(2) of such treaty in the English language.

9. Claims 43, 46, and 48-52 are rejected under 35 U.S.C. 102(b) as being anticipated by

Ghosh et al., Neuron, 15: 89-03, 1995. Independent claims 43 and 50 are drawn to nearly

identical methods in which two matched collections of cells are contacted with a candidate

modulator in the presence of an activator of an FGF signaling pathway or with an activator of an

FGF signaling pathway alone, and neural differentiation in the two cell collections is compared.

Claim 51 is drawn to apparently the same method for determining if the candidate modulator

modulates FGF-dependent neural differentiation; claim 52 recites contacting cells with the

candidate modulator in the presence of an activator of an FGF signaling pathway. In the

experiment shown in figure 9 in Ghosh et al., neural progenitor cells were incubated in the

presence of bFGF with or without the additional presence of either NT-3 (panel D) or anti-NT3

antibody (panel B). This experiment, therefore, teaches the instantly claimed methods, with NT-3

Application/Control Number: 10/789,266 Page 6

Art Unit: 1647

and anti-NT3 antibody as candidate modulators. NT-3 was observed to enhance neural differentiation (see panel D), thereby anticipating instant claim 48.

- 10. Claims 43-46 and 48-52 are rejected under 35 U.S.C. 102(b) as being anticipated by Lee et al., Nature Biotech. 18:675-678 (2000) (of record, IDS 01/18/2005). Lee et al. teach the generation of neurons from embryonic stem cells. In the experiment shown in figure 5 of Lee et al., embryonic stem cells (including neural progenitors derived from embryonic stem cells) were incubated in the presence of FGF with or without the additional presence of SHH. This experiment, therefore, teaches the instantly claimed methods, with SHH as a candidate modulator. SHH was observed to enhance neural differentiation (see panel B), thereby anticipating instant claim 48.
- 11. Claims 43-52 are rejected under 35 U.S.C. 102(e) as being anticipated by US Patent Application publication 20040092012 (Okano), filed October 3, 2001. Okano teaches methods for producing motor neurons from embryonic stem cells within embryoid bodies; the method employs a medium containing a fibroblast growth factor (FGF) and a sonic hedgehog protein [0004, 0019, 0020]. In the experiment shown in figures 7 and 8 of Okano, embryonic stem cells (including neural progenitors derived from embryonic stem cells) were incubated in the presence of FGF with or without the additional presence of noggin or SHH. These experiments, therefore, teach the instantly claimed methods, with noggin and SHH as candidate modulators, each of which were observed to enhance differentiation of motor neurons.

Conclusion

12. No claims are allowed.

Art Unit: 1647

Any inquiry concerning this communication or earlier communications from the examiner should be directed to Daniel C Gamett, Ph.D., whose telephone number is 571 272 1853. The examiner can normally be reached on M-F, 8:30-5:00.

If attempts to reach the examiner by telephone are unsuccessful, the examiner's supervisor, Brenda Brumback can be reached on 571 272 0961. The fax phone number for the organization where this application or proceeding is assigned is 571 273 8300.

Information regarding the status of an application may be obtained from the Patent Application Information Retrieval (PAIR) system. Status information for published applications may be obtained from either Private PAIR or Public PAIR. Status information for unpublished applications is available through Private PAIR only. For more information about the PAIR system, see http://pair-direct.uspto.gov. Should you have questions on access to the Private PAIR system, contact the Electronic Business Center (EBC) at 866-217-9197 (toll-free). If you would like assistance from a USPTO Customer Service Representative or access to the automated information system, call 800-786-9199 (IN USA OR CANADA) or 571-272-1000.

DCG

Art Unit 1647 20 April 2007 Jany GMWles

> GARY B. NICKOL, PH.D. SUPERVISORY PATENT EXAMINER TECHNOLOGY CENTER 1600